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	NEWS	1			Web Page for STN Seminar Schedule - N. America							
	NEWS		JAN	0.2	STN pricing information for 2008 now available							
	NEWS			16								
			01111		prophetic substances							
	NEWS	4	JAN	28	USPATFULL, USPAT2, and USPATOLD enhanced with new custom IPC display formats							
	NEWS	5	JAN	28	MARPAT searching enhanced							
	NEWS	6	JAN	28	USGENE now provides USPTO sequence data within 3 days of publication							
	NEWS	7	JAN	28	TOXCENTER enhanced with reloaded MEDLINE segment							
	NEWS			28	MEDLINE and LMEDLINE reloaded with enhancements							
	NEWS		FEB		STN Express, Version 8.3, now available							
	NEWS				PCI now available as a replacement to DPCI							
	NEWS				IFIREF reloaded with enhancements							
	NEWS				IMSPRODUCT reloaded with enhancements							
	NEWS				WPINDEX/WPIDS/WPIX enhanced with ECLA and current							
					U.S. National Patent Classification							
	NEWS	14	MAR	31	IFICDB, IFIPAT, and IFIUDB enhanced with new custom							
				-	IPC display formats							
	NEWS	1.5	MAR	3.1	CAS REGISTRY enhanced with additional experimental							
				-	spectra							
	NEWS	16	MAR	31	CA/CAplus and CASREACT patent number format for U.S.							
				-	applications updated							
	NEWS	17	MAR	31	LPCI now available as a replacement to LDPCI							
	NEWS				EMBASE, EMBAL, and LEMBASE reloaded with enhancements							
	NEWS				STN AnaVist, Version 1, to be discontinued							
	NEWS	20	APR	1.5	WPIDS, WPINDEX, and WPIX enhanced with new							
					predefined hit display formats							
	NEWS	21	APR	28	EMBASE Controlled Term thesaurus enhanced							
	NEWS				IMSRESEARCH reloaded with enhancements							
	NEWS	23	MAY	30	INPAFAMDB now available on STN for patent family							
					searching							
	NEWS	2.4	MAY	3.0	DGENE, PCTGEN, and USGENE enhanced with new homology							
					sequence search option							
	NEWS	25	JUN	06	EPFULL enhanced with 260,000 English abstracts							
	NEWS			06								
	NEWS	27	JUN	13	USPATFULL and USPAT2 updated with 11-character							
					patent numbers for U.S. applications							
	NEWS	28	JUN	19	CAS REGISTRY includes selected substances from web-based collections							
	MEMS	EVPI	PESS	FFRI	RUARY 08 CURRENT WINDOWS VERSION IS V8.3,							
	ишир	DAL KODO		AND CURRENT DISCOVER FILE IS DATED 20 FEBRUARY 2008								
				11111	CONTROL DISCOVERY I IN STREET TO FEBRUARY 2000							
	NEWS	HOU	2.5	STI	N Operating Hours Plus Help Desk Availability							
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STRUCTURE UPLOADED

=> dl1

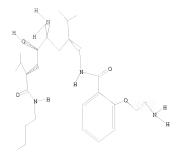
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=> d 11

L1 HAS NO ANSWERS

STR



Structure attributes must be viewed using STN Express guery preparation.

=> s 11 full

FULL SEARCH INITIATED 15:03:13 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 478 TO ITERATE

100.0% PROCESSED 478 ITERATIONS SEARCH TIME: 00.00.01 2 ANSWERS

L2 2 SEA SSS FUL L1

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FULL ESTIMATED COST 179.74 179.95

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=> s 12
L3
            2 L2
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=> d 13 1-2 ibib abs hitstr

L3 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:991333 CAPLUS

DOCUMENT NUMBER: 140:35967

TITLE: Methods of treating Alzheimers disease using aromatically substituted @-amino-alkanoic acid

amides and alkanoic acid diamides

INVENTOR(S): Maillard, Michel; Varghese, John PATENT ASSIGNEE(S): Elan Pharmaceuticals, Inc., USA SOURCE:

PCT Int. Appl., 286 pp. CODEN: PIXXD2

DOCUMENT TYPE: Patent. LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATEN	PATENT NO.					DATE										
					-											
WO 20	WO 2003103652					20031218		WO 2003-US18283					20030611			
W	: AE.	AG.	AL.	AM.	AT.	AU.	AZ.	BA.	BB.	BG,	BR.	BY.	BZ.	CA.	CH.	CN.
										EE,						
	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	LK,	LR,
	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	OM,	PH,
	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	TJ,	TM,	TN,	TR,	TT,	TZ,
	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW						
R	W: GH,	GM,	KΕ,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,	BY,
	KG,	KZ,	MD,	RU,	TJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,
	FI,	FR,	GB,	GR,	HU,	IE,	IT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,	TR,
	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG
AU 2003237546						2003	1222		AU 2	2003-	2375	46		2	0030	611
US 20060089355						2006	0060427			US 2005-517981				20050714		
PRIORITY APPLN. INFO.:									US 2	2002-	3877.	56P		P 2	0020	611
									WO 2	2003-	US18:	283		W 2	0030	611
OTHER SOURCE(S):					PAT	140:	3596	7								

OTHER SOURCE(S):

AB Disclosed are methods for treating Alzheimer's disease (no data), and other diseases (no data), and/or inhibiting beta-secretase enzyme (no data), and/or inhibiting deposition of amyloid β peptide (no data) in a mammal, using @-amino-alkanoic acid amides and alkanoic acid diamides (R1-X1-NH-X2-CH(R2)CH2CH(R3)CH(OH)CH2CH(R4)C(O)NH-R5 (I); variables defined below; e.g. (2S, 4S, 5S, 7R)-N-(4-amino-7-butyl-7-carbamoyl-

5-hydroxy-2-isopropyloctyl)-3-methoxy-2-(3-methoxypropoxy)benzamide). Many example prepns. are included but all of them comprise an English translation of a German patent (EP 0716077 A1; 1996; CA file accession number 125:167576). For I: R1 is a 2-RA-3-RB-Ph radical, a 2-RA-4-Rc-Ph radical, a 2-RA-pyridin-3-vl radical a 3-RA-pyridin-2-vl radical or a 1-RD-indol-3-vl radical, wherein one of the radicals RA and RB is an

aliphatic or heterocycloaliph.-aliphatic radical or free or aliphatically, araliphatically or heteroaraliphatically etherified hydroxy and the other is H, an aliphatic radical or free or esterified or amidated carboxy, RC is H, an aliphatic radical, free or aliphatically, araliphatically,

heteroaraliphatically or heteroarylaliphatically etherified hydroxy or an unsubstituted or heteroaliphatically substituted amino group, and RD is an aliphatic, araliph. or heteroaliph. radical, one of the radicals X1 and X2 is carbonyl and the other is methylene, R2 is an aliphatic radical, R3 is

unsubstituted or aliphatically substituted amino, R4 is an aliphatic or araliph. radical, and R5 is an aliphatic or cycloaliph.—aliphatic radical or an optionally hydrogenated and/or oxo—substituted heteroaryl radical or an optionally hydrogenated and/or oxo—substituted heteroaryl or heteroaliph. radical bonded via a C atom.

T 179995-26-7P 180183-70-4P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; methods of treating Alzheimers disease using aromatically substituted ω -amino-alkanoic acid amides and alkanoic acid diamides)

RN 179995-26-7 CAPLUS

CN Benzamide, N-[(28,48,58,78)-4-amino-7-[(butylamino)carbonyl]-5-hydroxy-8-methyl-2-(1-methylethyl)nonyl]-2-(2-aminoethoxy)-, dihydrochloride (9CI) (CA INDEX NAMB)

Absolute stereochemistry.

● 2 HC1

RN 180183-70-4 CAPLUS

CN Benzamide, N-[(2S,4S,5S,7S)-4-amino-7-[(butylamino)carbonyl]-5-hydroxy-8-methyl-2-(1-methylethyl)nonyl]-2-(2-aminoethoxy)- (CA INDEX NAME)

Absolute stereochemistry.

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1996:501390 CAPLUS DOCUMENT NUMBER: 125:167576

ORIGINAL REFERENCE NO.: 125:31396h,31397a
TITLE: Aryl-substituted @-aminoalkanamides and diamides

and their use as renin inhibitors

INVENTOR(S): Maibaum, Juergen K.; Rigollier, Pascal; Herold, Peter; Cohen, Nissim C.; Goeschke, Richard; Stutz, Stefan

PATENT ASSIGNEE(S): Ciba-Geigy A.-G., Switz.

SOURCE:

Eur. Pat. Appl., 90 pp.

CODEN: EPXXDW

DOCUMENT TYPE: LANGUAGE:

Patent German

FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE			
EP 716077	A1	19960612	EP 1995-810743	19951129			
R: AT, BE, CH,	DE, DK	, ES, FR,	GB, GR, IE, IT, LI, LU,	NL, PT, SE			
FI 9505836	A	19960609	FI 1995-5836	19951204			
CA 2164571	A1	19960609	CA 1995-2164571	19951206			
ZA 9510354	A	19960610	ZA 1995-10354	19951206			
AU 9540266	A	19960613	AU 1995-40266	19951206			
US 5641778	A	19970624	US 1995-568332	19951206			
NO 9504975	A	19960610	NO 1995-4975	19951207			
JP 08231485	A	19960910	JP 1995-319220	19951207			
CN 1136556	A	19961127	CN 1995-113102	19951207			
HU 74454	A2	19961230	HU 1995-3508	19951207			
PRIORITY APPLN. INFO.:			CH 1994-3724	A 19941208			
OTHER SOURCE(S):	MARPAT	125:1675	76				
GI							

The N-[amino(hydroxy)oxooctyl]amides I (R1 = aryl; R2 = aliphatic group; R3 = AB aminoalkyl group; R5 = alkyl, cycloalkyl, etc.; X1, X2 = methylene, carbonyl) were disclosed. I are useful as renin inhibitors and for the treatment of hypertension. A claimed example compound is (2S, 4S, 5S, 7R)-N-(4-amino-7-butyl-7-carbamoyl-5-hydroxy-2-isopropyloctyl)-3methoxy-2-(3-methoxypropoxy)benzamide (II).

II

179995-26-7P 180183-70-4P RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of N-[amino(hydroxy)oxooctyl]amides as renin inhibitors) RN 179995-26-7 CAPLUS

CN Benzamide, N-[(2S,4S,5S,7S)-4-amino-7-[(butylamino)carbony1]-5-hydroxy-8methyl-2-(1-methylethyl)nonyl]-2-(2-aminoethoxy)-, dihydrochloride (9CI) (CA INDEX NAME)

Absolute stereochemistry.

●2 HC1

RN 180183-70-4 CAPLUS

CN Benzamide, N-[(2S,4S,5S,7S)-4-amino-7-[(butylamino)carbonyl]-5-hydroxy-8-methyl-2-(1-methylethyl)nonyl]-2-(2-aminoethoxy)- (CA INDEX NAME)

Absolute stereochemistry.

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Connection closed by remote host